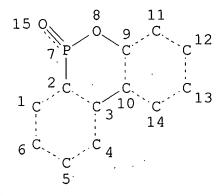
FILE 'HCAPLUS' ENTERED AT 11:10:24 ON 24 APR 2006
L8 14 S L4(L)RACT+NT/RL AND L7(L)PREP+NT/RL

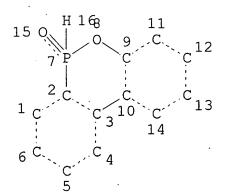
FILE 'REGISTRY' ENTERED AT 11:12:36 ON 24 APR 2006

FILE 'REGISTRY' ENTERED AT 11:12:55 ON 24 APR 2006

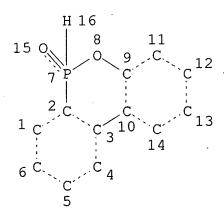
=> str l1 :dis



:att 7 h vn se
:dis



:mov 16 u2, dis



:end

L9 STRUCTURE CREATED

=> s 19 sub=14 sam

SAMPLE SUBSET SEARCH INITIATED 11:14:02 FILE 'REGISTRY'

SAMPLE SUBSET SCREEN SEARCH COMPLETED - 99 TO ITERATE

100.0% PROCESSED

99 ITERATIONS

9

ANSWERS

SEARCH TIME: 00.00.01

PROJECTIONS (WITHIN SPECIFIED SUBSET): ONLINE **COMPLETE**

PROJECTED ITERATIONS (WITHIN SPECIFIED SUBSET): 1384 TO

2576

PROJECTED ANSWERS (WITHIN SPECIFIED SUBSET): 9 TO

360

L10 9 SEA SUB=L4 SSS SAM L9

=> s 19 sub=14 full

FULL SUBSET SEARCH INITIATED 11:14:10 FILE 'REGISTRY'
FULL SUBSET SCREEN SEARCH COMPLETED - 1846 TO ITERATE

100.0% PROCESSED 1846 ITERATIONS 127

ANSWERS

SEARCH TIME: 00.00.01

L11 127 SEA SUB=L4 SSS FUL L9

=> b hcap

storing

COST IN U.S. DOLLARS SINCE FILE TOTAL

FULL ESTIMATED COST ENTRY SESSION 40.28 645.29

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SINCE FILE TOTAL ENTRY SESSION

CA SUBSCRIBER PRICE 0.00 -10.50

FILE 'HCAPLUS' ENTERED AT 11:14:15 ON 24 APR 2006 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS.

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FILE COVERS 1907 - 24 Apr 2006 VOL 144 ISS 18 FILE LAST UPDATED: 23 Apr 2006 (20060423/ED)

New CAS Information Use Policies, enter HELP USAGETERMS for details.

This file contains CAS Registry Numbers for easy and accurate substance identification.

 \Rightarrow 111 (l) ract+nt/rl and 17(l) prep+nt/rl

L11 IS NOT A RECOGNIZED COMMAND

The previous command name entered was not recognized by the system. For a list of commands available to you in the current file, enter "HELP COMMANDS" at an arrow prompt (=>).

 \Rightarrow s l11 (l) ract+nt/rl and l7(l) prep+nt/rl

582 L11

2845537 RACT+NT/RL (10 TERMS)

168 L11 (L) RACT+NT/RL

165 L7

3456277 PREP+NT/RL (18 TERMS)

61 L7(L) PREP+NT/RL

L12 10 L11 (L) RACT+NT/RL AND L7(L) PREP+NT/RL

=> d ibib abs hitstr 112 tot

L12 ANSWER 1 OF 10 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2006:293401 HCAPLUS

DOCUMENT NUMBER: 144:332992

TITLE: Fireproofing finishes containing phosphonic

acid

esters and manufacture of fire-resistant

fibers using

them

INVENTOR(S): Kobayashi, Junichi; Ishikawa, Akira;

Kanehira, Ryoji

PATENT ASSIGNEE(S): Marubishi Oil Chemical Co., Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 20 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO. DATE	KIND	DATE	APPLICATION NO.
JP 2006083491 ·	A2	20060330	JP 2004-270028
20040916			

```
20040916
     The finishes contain RC6H4OQ [I; R = H, (un) substituted
hydrocarbyl; Q =
     9,10-dihydro-10-oxo-9-oxa-10-phosphaphenanthren-10-yl] and are
contacted
     to fibers to give the fire-resistant fibers. Manufacture of I
by reaction of
     9,10-dihydro-9-oxa-10-phosphaphenanthrene 10-oxide (II) with
     compds. in the presence of amines and dehydrohalogenation
reaction of the
     resulting XQ (X = halo) with RC6H4OH is also claimed.
    was added dropwise to CH2Cl2 containing II 32.4, phenol 14.1,
and Et3N 17.2 g
     at \leq 15^{\circ} and stirred for 1 h to give 43.5 g I (R = H), 40
     parts of which was added to a mixture of polyoxyethylene
distyrenated phenol
     ether sulfate 5, 10% aqueous CM-cellulose solution 2, and H2O 53
parts and
     dispersed to give a fireproofing finish. A 90/10 regular
     polyester/cationic dyeable polyester fabric was padded with a
liquid containing
     20% of the finish, dried, heat-set, washed with soda ash, and
dried to
     show good fire resistance even after washing or dry cleaning.
     36240-30-9P 55217-59-9P, 6-Phenoxy-6H-
IT
     Dibenz[c,e][1,2]oxaphosphorin-6-oxide 880138-78-3P,
     6-(3-Methylphenoxy)-6H-Dibenz[c,e][1,2]oxaphosphorin-6-oxide
     880138-79-4P, 6-(4-tert-Butylphenoxy)-6H-
     Dibenz[c,e][1,2]oxaphosphorin-6-oxide 880138-80-7P,
     6-(4-n-Octylphenoxy)-6H-Dibenz[c,e][1,2]oxaphosphorin-6-oxide
     RL: IMF (Industrial manufacture); MOA (Modifier or additive
     use); PRP (Properties); PREP (Preparation); USES (Uses)
        (fireproofing finishes based on phosphonic acid esters with
good
        washfastness)
RN
     36240-30-9 HCAPLUS
     6H-Dibenz[c,e][1,2] oxaphosphorin, 6-([1,1'-biphenyl]-2-yloxy)-,
6-oxide
     (9CI)
            (CA INDEX NAME)
```

PRIORITY APPLN. INFO.:

JP 2004-270028

RN 55217-59-9 HCAPLUS
CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 6-phenoxy-, 6-oxide (9CI) (CA
INDEX
NAME)

RN 880138-78-3 HCAPLUS CN 6H-Dibenzo[c,e][1,2]oxaphosphorin, 6-(3-methylphenoxy)-, 6-oxide (9CI) (CA INDEX NAME)

RN 880138-79-4 HCAPLUS
CN 6H-Dibenzo[c,e][1,2]oxaphosphorin,
6-[4-(1,1-dimethylethyl)phenoxy]-,
6-oxide (9CI) (CA INDEX NAME)

RN 880138-80-7 HCAPLUS

CN 6H-Dibenzo[c,e][1,2]oxaphosphorin, 6-(4-octylphenoxy)-, 6-oxide

(9CI) (CA

INDEX NAME)

IT 35948-25-5, 9,10-Dihydro-9-oxa-10-phosphaphenanthrene 10-oxide

RL: RCT (Reactant); RACT (Reactant or reagent)

(fireproofing finishes based on phosphonic acid esters with

good

washfastness)

RN 35948-25-5 HCAPLUS

CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 6-oxide (9CI) (CA INDEX NAME)

L12 ANSWER 2 OF 10 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2005:141621 HCAPLUS

DOCUMENT NUMBER: 143:172925
TITLE: Arylation of

6H-dibenzo[c,e][1,2 λ 5]oxaphosphini

ne 6-oxide

AUTHOR(S):

Beletskaya, I. P.; Neganova, E. G.; Veits,

Yu. A.

CORPORATE SOURCE:

Faculty of Chemistry, Lomonosov Moscow State

University, Moscow, 119992, Russia

SOURCE:

Russian Journal of Organic Chemistry (2004),

40(12),

1782-1786

CODEN: RJOCEQ; ISSN: 1070-4280

PUBLISHER:

MAIK Nauka/Interperiodica Publishing

DOCUMENT TYPE:

Journal

LANGUAGE:

English

OTHER SOURCE(S):

CASREACT 143:172925

AB Arylation and alkylation of 6H-dibenzo[c,e][1,2 λ 5]oxaphosphinine 6-oxide at the phosphorus atom was accomplished.

Tetrafluoro-4-pyridyl

fragment was introduced via reaction of 6-trimethylsiloxy-6H-dibenzo[c,e][1,2]oxaphosphinine with pentafluoropyridine. The arylation

of the title compound with aryl iodides containing both electron-acceptor and

electron-donor substituents was effected under catalysis by palladium or

nickel complexes.

IT 35948-25-5

RL: RCT (Reactant); RACT (Reactant or reagent)

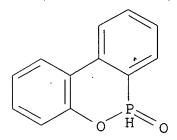
(palladium or nickel catalyzed arylation of

dibenzooxaphosphinine

oxide)

RN 35948-25-5 HCAPLUS

CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 6-oxide (9CI) (CA INDEX NAME)



IT 861105-63-7P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP

(Preparation); RACT (Reactant or reagent)

(palladium or nickel catalyzed arylation of

dibenzooxaphosphinine

oxide)

RN 861105-63-7 HCAPLUS

CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 6-[(trimethylsilyl)oxy]- (9CI) (CA

REFERENCE COUNT:

25 THERE ARE 25 CITED REFERENCES

AVAILABLE FOR THIS

RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

L12 ANSWER 3 OF 10 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2005:140870 HCAPLUS

DOCUMENT NUMBER:

142:198207

TITLE:

Process for the preparation of

9,10-dihydro-9-oxa-10-

organophosphaphenanthrene 10-oxide and

derivatives of

INVENTOR (S):

the same substituted on the phenyl groups

Dittrich, Uwe; Just, Berthold; Doring,

Manfred;

Ciesielski, Michael

PATENT ASSIGNEE(S):

Germany

SOURCE:

U.S. Pat. Appl. Publ., 12 pp.

CODEN: USXXCO

DOCUMENT TYPE:

Patent -

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

1

PATENT INFORMATION:

	PATENT NO.		KINI) _.	DATE			APPL	ICAT:	ION I	. O <i>v</i>		,
DATE				-								-	٠
	US 2005038279		A1		2005	0217		US. 2	004-	9188	38		
2004	DE 10338116		A1		2005	0317		DE 2	003-	1033	8116		
2003	0815 EP 1512690		A1		2005	0309		EP 2	004-	1882	9 .		(
2004	0809 R: AT, BE,	CH,	DE,	DK,	, ĘS,	FR,	GB,	GR,	ΙŤ,	LI,	LU,	NL,	SE,
MC,	PT, IE, SI,	LT,	LV,	FΙ	, RO,	MK,	CY,	AL,	TR,	BG,	CZ,	EE,	HU,
•	SK, HR RITY APPLN. INFO. 0815	. :						DE 2	003-	1033	8116	j	A

OTHER SOURCE(S): CASREACT 142:198207; MARPAT 142:198207

AB A process is provided for the preparation of

9,10-dihydro-9-oxa-10-

organophosphaphenanthrene 10-oxide and derivs. of same substituted on the

Ph groups, in which: (a)

9,10-dihydro-9-oxa-10-phosphaphenanthrene

. 10-oxide (DOP) or a derivative of same is reacted in the presence of at least

one mono- or polyhydric alc. with at least one ortho ester with formation

of a first intermediate product, (b) the intermediate product from step

(a) is optionally reacted with at least one further mono- or polyhydric

alc. with formation of a further intermediate product and (c) the intermediate product from steps (a) or (b) is transformed by addition of

catalytic quantities of alkylation agent into 9,10-dihydro-9-oxa-10-

organophosphaphenanthrene 10-oxide or a derivative of same substituted on the

Ph groups.

IT 35948-25-5, 9,10-Dihydro-9-oxa-10-phosphaphenanthrene-10-oxide

RL: RCT (Reactant); RACT (Reactant or reagent)
(preparation of substituted Ph group derivs. of dihydro
oxaorganophosphaphenanthrene oxide useful as flame retardants)

RN 35948-25-5 HCAPLUS

CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 6-oxide (9CI) (CA INDEX NAME)

IT 37632-28-3P 194091-96-8P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation of substituted Ph group derivs. of dihydro oxaorganophosphaphenanthrene oxide useful as flame retardants) 37632-28-3 HCAPLUS

RN 37632-28-3 HCAPLUS
CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 6-methoxy- (9CI) (CA INDEX NAME)

RN 194091-96-8 HCAPLUS

CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 6-ethoxy- (9CI) (CA INDEX NAME)

IT - 585573-01-9P

RL: SPN (Synthetic preparation); PREP (Preparation)

(preparation of substituted Ph group derivs. of dihydro oxaorganophosphaphenanthrene oxide useful as flame retardants)

RN 585573-01-9 HCAPLUS

CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 6-(1-methylethoxy)- (9CI) (CA

INDEX

NAME)

L12 ANSWER 4 OF 10 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2005:135413 HCAPLUS

DOCUMENT NUMBER: 142:198206

TITLE: Process for the preparation of

9,10-dihydro-9-oxa-10-

organophosphaphenanthrene 10-oxide and

derivatives of

the same substituted on the phenyl groups Dittrich, Uwe; Just, Berthold; Doering, INVENTOR(S): Manfred; Ciesielski, Michael Schill & Seilacher "struktol" PATENT ASSIGNEE(S): Aktiengesellschaft, Germany Eur. Pat. Appl., 19 pp. SOURCE: CODEN: EPXXDW DOCUMENT TYPE: Patent LANGUAGE: German FAMILY ACC. NUM. COUNT: PATENT INFORMATION: KIND DATE APPLICATION NO. PATENT NO. DATE A1 20050216 EP 2004-18830 EP 1506968 20040809 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK, HR A1 20050317 DE 2003-10338131 DE 10338131 20030815 20040813 DE 2003-10338131 A PRIORITY APPLN. INFO.: 20030815 OTHER SOURCE(S): CASREACT 142:198206; MARPAT 142:198206 A process is provided for the preparation of 9,10-dihydro-9-oxa-10organophosphaphenanthrene 10-oxide and derivs. of same substituted on the Ph groups, in which: (a) 9,10-dihydro-9-oxa-10-phosphaphenanthrene 10-oxide (DOP) or a derivative of same is reacted in the presence of at least one mono- or polyhydric alc. with at least one ortho ester with formation of a first intermediate product, (b) the intermediate product from step (a) is optionally reacted with at least one further mono- or alc. with formation of a further intermediate product and (c) the intermediate product from steps (a) or (b) is transformed by addition of catalytic quantities of alkylation agent into 9,10-dihydro-9-oxa-10organophosphaphenanthrene 10-oxide or a derivative of same

substituted on the

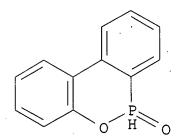
Ph groups.

IT 35948-25-5, 9,10-Dihydro-9-oxa-10-phosphaphenanthrene-10-oxide RL: RCT (Reactant); RACT (Reactant or reagent)

(preparation of substituted Ph group derivs. of dihydro oxaorganophosphaphenanthrene oxide useful as flame retardants)

RN 35948-25-5 HCAPLUS

CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 6-oxide (9CI) (CA INDEX NAME)



IT 37632-28-3P

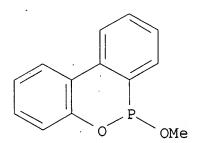
RL: RCT (Reactant); SPN (Synthetic preparation); PREP

(Preparation); RACT (Reactant or reagent)

(preparation of substituted Ph group derivs. of dihydro oxaorganophosphaphenanthrene oxide useful as flame retardants)

RN 37632-28-3 HCAPLUS

CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 6-methoxy- (9CI) (CA INDEX NAME)



IT 103764-64-3P 194091-96-8P 585573-01-9P

RL: SPN (Synthetic preparation); PREP (Preparation)

(preparation of substituted Ph group derivs. of dihydro oxaorganophosphaphenanthrene oxide useful as flame retardants)

RN 103764-64-3 HCAPLUS

CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 6-methoxy-, 6-oxide (9CI) (CA INDEX

NAME)

RN 194091-96-8 HCAPLUS

CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 6-ethoxy- (9CI) (CA INDEX NAME)

RN 585573-01-9 HCAPLUS

CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 6-(1-methylethoxy)- (9CI) (CA

INDEX

NAME)

REFERENCE COUNT:

5 THERE ARE 5 CITED REFERENCES AVAILABLE

FOR THIS

RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

L12 ANSWER 5 OF 10 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2003:678817 HCAPLUS

DOCUMENT NUMBER: 139:197621

TITLE: Method for producing

6-alkoxy-(6H)-dibenz[c,e][1,2]-

oxaphosphorins

INVENTOR(S):

Sprenger, Stephan; Ciesielski, Michael;

Kollann,

Carsten; Doering, Manfred

PATENT ASSIGNEE(S):

Forschungszentrum Karlsruhe G.m.b.H., Germany

SOURCE:

PCT Int. Appl., 24 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

German

FAMILY ACC. NUM. COUNT:

1 .

PATENT INFORMATION:

PATENT NO.

KIND DATE

APPLICATION NO.

DATE

WO 2003070736 A1 20030828

WO 2003-EP1368

20030212

W: JP, US

RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR,

HU, IE,

IT, LU, MC, NL, PT, SE, SI, SK, TR

DE 10206982 A1 20030904 DE 2002-10206982

20020220

DE 10206982

B4 20040325

EP 1476453

A1 20041117 EP 2003-702624

20030212

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE,

MC, PT,

IE, SI, FI, CY, TR, BG, CZ, EE, HU, SK

JP 2005517740 T2 20050616 JP 2003-569643

20030212

US 2005176983 A1 20050811 US 2003-505201

20030212

PRIORITY APPLN. INFO.:

DE 2002-10206982

Α

20020220

WO 2003-EP1368 W

20030212

OTHER SOURCE(S):

CASREACT 139:197621; MARPAT 139:197621

GΙ

AB The invention relates to a method for producing 6-alkoxy-(6H)-dibenz[c,e][1,2]-oxaphosphorins, whereby

6H-dibenz[c,e][1,2]-oxaphosphorin-

6-oxides I (R3, R4 = alkyl, alkoxy, alkylthio, alkenyl, alkynyl, aryl,

heteroaryl, cycloalkyl) are used as adduct. Thus, reaction of 6H-dibenz[c,e][1,2]-oxaphosphorin 6-oxide with HCl in methanol at 85° for 45 min followed by treatment with concentrate HCl for 5

h and

tri-Me orthoformate for 30 min gave 87%
6-methoxy-6H-dibenz[c,e][1,2]-

oxaphosphorin.

IT 35948-25-5, 6H-Dibenz[c,e][1,2]-oxaphosphorin 6-oxide

RL: RCT (Reactant); RACT (Reactant or reagent)
(method for producing alkoxydibenzoxaphosphorins starting from dibenzoxaphosphorin oxides)

RN 35948-25-5 HCAPLUS

CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 6-oxide (9CI) (CA INDEX NAME)

37632-28-3P, 6-Methoxy-6H-dibenz[c,e][1,2]-oxaphosphorin 194091-96-8P, 6-Ethoxy-6H-dibenz[c,e][1,2]-oxaphosphorin 585573-01-9P, 6-Isopropoxy-6H-dibenz[c,e][1,2]-oxaphosphorin RL: SPN (Synthetic preparation); PREP (Preparation)

(method for producing alkoxydibenzoxaphosphorins starting from dibenzoxaphosphorin oxides)

RN 37632-28-3 HCAPLUS

CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 6-methoxy- (9CI) (CA INDEX NAME)

RN 194091-96-8 HCAPLUS CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 6-ethoxy- (9CI) (CA INDEX NAME)

RN 585573-01-9 HCAPLUS
CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 6-(1-methylethoxy)- (9CI) (CA
INDEX
NAME)

REFERENCE COUNT:

3 THERE ARE 3 CITED REFERENCES AVAILABLE

FOR THIS .

RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

L12 ANSWER 6 OF 10 HCAPLUS COPYRIGHT 2006 ACS on STN ACCESSION NUMBER: 2003:274797 HCAPLUS

DOCUMENT NUMBER:

138:288481

TITLE:

Phosphorus-containing fire-resistant curing

agents and

epoxy resins, advanced epoxy resins, and

cured epoxy

resins containing them

Wang, Chun Shan; Hsieh, Cheng Yueh; Lin, INVENTOR(S):

Ching Yuan

Taiwan

PATENT ASSIGNEE(S): SOURCE:

Jpn. Kokai Tokkyo Koho, 29 pp.

CODEN: JKXXAF

DOCUMENT TYPE:

Patent

LANGUAGE:

Japanese

FAMILY ACC. NUM. COUNT:

1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.
DATE			·
JP 2003105058	A2	20030409	JP 2001-386377
20011219			
JP 3653247	B2	20050525	
TW 593526	В	20040621	TW 2001-90123251
20010920		,	
US 2003120021	A1	20030626	US 2002-66455
20020130			
US 6797821	B2	20040928	
US 2005004339	A1	20050106	US 2004-896567
20040722			
PRIORITY APPLN. INFO.:			TW 2001-90123251 A
20010920			
			US 2002-66455 A3
20020130			
GI			

$$HO \longrightarrow X \longrightarrow OH$$

$$(Q) j(H)2-jN$$
 N
 $N(H) 2-j(Q) j$ II

AB The curing agents are selected from I, NH2-iQiC6H4-p-XC6H4-p-NH2-jQj,

triazine derivs. II, N.tplbond.CN:C(NH2-jQj)NH2-iQi, O'C(NH2)2NHC(O'):NH,

H2NC(:NH)NHC(Q'):NH, and N.tplbond.CNH1-kQ'kC(:NQ')NH2-iQ'i [1, m. i. i =

0-2; 1 + m > 0; 0 < i + j < 4; k = 0-1; i + k < 3; Z = NH2, Me, Ph; X =

direct link, CH2, CMe2, cyclohexylidene, O, S, SO2; Q = Q'CR1R2,
O': O' =

6-oxido-6H-dibenz[c,e][1,2]oxaphósphorin-6-yl, Ar2P(O); R1, R2 = H, C1-18

alkyl, C6-18 (un) substituted aryl, C6-18 (un) substituted arylmethylene; Ar

= C1-4 alkyl- or C6-18 aryl-(un)substituted Ph or phenoxy]. Epoxy resins

containing the curing agents are useful for semiconductor device packaging.

Thus, bisphenol A was reacted with equimolar (6-oxido-6H-dibenz[c,e][1,2]oxaphosphorin-6-yl)methanol in the presence of AcOK to

aive

[(6-oxido-6H-dibenz[c,e][1,2]oxaphosphorin-6-yl)methyl]bisphenol A, 228 g of which was treated with 564 g bisphenol A diglycidyl ether at

160° for 2 h in the presence of EtPPh3Cl to give an epoxy resin. The epoxy resin was cured with a novolak to show 5% weight loss temperature

387° in air and N and good fire resistance.

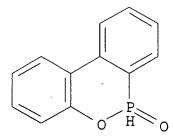
IT 507264-76-8P 507264-78-0P

RL: IMF (Industrial manufacture); RCT (Reactant); PREP (Preparation); RACT (Reactant or reagent)

(phosphorus-containing fire-resistant curing agents for epoxy resins)

RN 507264-76-8 HCAPLUS
CN 1,3,5-Triazine-2,4,6-triamine,
N-(6-oxido-6H-dibenz[c,e][1,2]oxaphosphorin-6-yl)-(9CI) (CA INDEX NAME)

ONE OR MORE TAUTOMERIC DOUBLE BONDS NOT DISPLAYED IN THE STRUCTURE RN 507264-78-0 HCAPLUS CN Guanidine, N-cyano-N'-(6-oxido-6H-dibenz[c,e][1,2]oxaphosphorin-6-yl)-(9CI) (CA INDEX NAME)



HCAPLUS COPYRIGHT 2006 ACS on STN L12 ANSWER 7 OF 10

2003:134074 HCAPLUS ACCESSION NUMBER:

DOCUMENT NUMBER: 138:321381

The role of ligand transformations on the TITLE:

performance

of phosphite- and phosphinite-based palladium

catalysts in the Suzuki reaction

Bedford, Robin B.; Hazelwood, Samantha L.; AUTHOR(S):

Limmert,

Michael E.; Brown, John M.; Ramdeehul,

Shailesh:

Cowley, Andrew R.; Coles, Simon J.;

Hursthouse,

Michael B.

CORPORATE SOURCE:

School of Chemistry, University of Exeter,

Exeter, EX4

40D, UK

SOURCE:

Organometallics (2003), 22(7), 1364-1371

CODEN: ORGND7; ISSN: 0276-7333

American Chemical Society PUBLISHER:

Journal DOCUMENT TYPE:

English LANGUAGE:

CASREACT 138:321381 OTHER SOURCE(S):

The ortho-metalated complex [$\{Pd(\mu-C1)\}\{\kappa P, \kappa C-P(OC6H2-2, 4-1)\}\}$ tBu2) (OC6H3-2,4-tBu2)2}}2] reacts with phenylboronic acid hydrate and

K2CO3 in dimethylacetamide to give oxo-bridged diaryl phosphite complex

 $[Pd(\kappa P, \kappa C, \kappa O - \mu 2 - O - P(O) (OC6H2 - 2, 4 - tBu2) (OC6H3 - 2, 4 - tBu2)]$

tBu2)(DMAc)}] (11). When the reaction is repeated in DMF, the coupling

product, 3,3',5,5'-tetra-tert-butyl-2,2'-biphenol (12) was isolated. The

reaction of palladium dichloride with phosphinite PiPr2(OC6H4-4-Et) in

2-methoxyethanol followed by recrystn. in the presence of ethanol gave the

palladium complex of the transesterified phosphinite ligand, trans-[PdCl2{PiPr2(OEt)}2] (14). The mol. structure of 11, 12 and 14 was

confirmed by x-ray crystallog. To determine whether related solvolytic

processes have an effect on catalytic activity, the performance of a range

of catalysts with "hydrolyzed" and "nonhydrolyzed" ligands was assessed in

the Suzuki coupling of aryl bromides. Palladium ortho-metalated dimethylbenzylamine and phosphite complexes with extra hydroxyphosphinite

and secondary phosphite ligands, [Pd(C6H4CH2NMe2- κ C, κ N)(L1- κ P)] (16, L1 = 6-hydroxy-6H-dibenzo[c,e][1,2]-oxaphosphorin), [Pd(L2- κ C, κ P)Cl]2 (3e, L2 = (2,4-di-tert-

butylphenyl) (methylenebis-2,2'-[6-tert-butyl-4-methylphenyl])
phosphite),

and in situ formed [Pd(C6H4CH2NMe2- κ C, κ N)(L2- κ P)] and [Pd(C6H4CH2NMe2- κ C, κ N)(L3- κ P)] (L3 =

(hydroxy) (methylenebis-2,2'-[6-tert-butyl-4-methylphenyl])
phosphite) were

tested as Suzuki coupling catalysts, showing moderate activity. In some

cases it was evident that hydrolysis plays a significant role on the

catalytic activity; however, this depends not only on the ligand, but also

on the combination of ligand and palladium precursor.

IT 512778-81-3P

RL: CAT (Catalyst use); SPN (Synthetic preparation); PREP (Preparation); USES (Uses)

(Suzuki coupling catalyst; preparation and Suzuki coupling catalytic

activity of palladium cyclometalated complexes with partially hydrolyzed ligands)

RN 512778-81-3 HCAPLUS

CN Palladium, [2-[(dimethylamino-κN)methyl]phenyl-κC](6-hydroxy-6H-dibenz[c,e][1,2]oxaphosphorin-κP6)(trifluoroacetato-κO)-, (SP-4-3)- (9CI) (CA INDEX NAME)

IT 35948-25-5

RL: CAT (Catalyst use); RCT (Reactant); RACT (Reactant or reagent); USES (Uses)

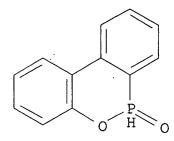
(Suzuki coupling co-catalyst, complexation; preparation and Suzuki coupling

catalytic activity of palladium cyclometalated complexes with partially

hydrolyzed ligands)

35948-25-5 HCAPLUS RN

6H-Dibenz[c,e][1,2]oxaphosphorin, 6-oxide (9CI) (CA INDEX NAME) CN



THERE ARE 35 CITED REFERENCES REFERENCE COUNT: 35

AVAILABLE FOR THIS

RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

HCAPLUS COPYRIGHT 2006 ACS on STN L12 ANSWER 8 OF 10

ACCESSION NUMBER:

2001:371569 HCAPLUS

DOCUMENT NUMBER:

134:354012

TITLE:

Synthesis of organophosphorus compounds and

their

metal salts.

INVENTOR(S):

Saito, Toranosuke; Ikemoto, Kenichi; Horii,

Hisashi

PATENT ASSIGNEE(S):

Sanko Kaihatsu Kagaku Kenkyusho K. K.,

Japan; Saito

Kaseihin Kenkyusho Y. K.

SOURCE:

Jpn. Kokai Tokkyo Koho, 8 pp.

CODEN: JKXXAF

DOCUMENT TYPE:

Patent

LANGUAGE:

Japanese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

DATE	PATENT NO.	KIND	DATE	APPLICATION NO.
19991	RITY APPLN. INFO.:	A2	20010522	JP 1999-322773 JP 1999-322773

OTHER SOURCE(S):

MARPAT 134:354012

GΪ

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AB Cyclic organophosphorus compound I, useful as fire retardant and stabilizer

for polymeric electronic and optical materials, is synthesized by hydrogen

peroxide oxidation of compound II or III in the presence of water and in the

presence or absence of an inert polar organic solvent followed by dehydrocyclization (X1-3 = H, halogen, alkyl, cycloalkyl, aryl, aralkyl).

IT 36240-31-0P 69151-14-0P 121166-84-5P

RL: IMF (Industrial manufacture); PREP (Preparation)

(synthesis of organophosphorus compds. and their metal salts) 36240-31-0 HCAPLUS

CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 6-hydroxy-, 6-oxide (9CI) (CA INDEX

NAME)

RN 69151-14-0 HCAPLUS

CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 6-hydroxy-, 6-oxide, zinc salt (9CI)

(CA INDEX NAME)

●1/2 Zn

RN 121166-84-5 HCAPLUS

CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 6-hydroxy-, 6-oxide, aluminum salt (9CI)

(CA INDEX NAME)

●1/3 Al .

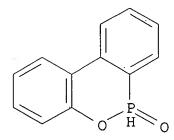
IT 35948-25-5, HCA

RL: RCT (Reactant); RACT (Reactant or reagent)

(synthesis of organophosphorus compds. and their metal salts)

RN 35948-25-5 HCAPLUS

CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 6-oxide (9CI) (CA INDEX NAME)



L12 ANSWER 9 OF 10 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1998:433491 HCAPLUS

DOCUMENT NUMBER: 129:149025

TITLE: Synthesis and Characterization of Novel

6-Substituted

4-Phenyl-6H-dibenz[c,e][1,2]oxaphosphorins

AUTHOR(S): Qureshi, Asfia; Hay, Allan S.

CORPORATE SOURCE: Department of Chemistry, McGill University,

Montreal,

QC, H3A 2K6, Can.

SOURCE: Journal of Chemical Research, Synopses

(1998), (7),

355, 1601-1615

CODEN: JRPSDC; ISSN: 0308-2342

PUBLISHER: Royal Society of Chemistry

DOCUMENT TYPE: Journal LANGUAGE: English

AB Novel 6-substituted 4-phenyl-6H-dibenz[c,e][1,2]oxaphosphorins

were

synthesized, starting from the reaction of 2,6-diphenylphenol

with P

trichloride using Zn chloride as catalyst.

IT 67362-63-4P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP

(Preparation); RACT (Reactant or reagent)

(preparation of dibenzoxaphosphorins from phenols and phosphorus

trichloride)

RN 67362-63-4 HCAPLUS

CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 4-phenyl-, 6-oxide (9CI) (CA INDEX

.NDEA

NAME)

IT 210899-90-4P 210899-99-3P

RL: SPN (Synthetic preparation); PREP (Preparation) (preparation of dibenzoxaphosphorins from phenols and phosphorus

trichloride)

RN 210899-90-4 HCAPLUS

CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 6-phenoxy-4-phenyl- (9CI) (CA INDEX

NAME)

RN 210899-99-3 HCAPLUS CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 6-phenoxy-4-phenyl-, 6-oxide (9CI) (CA

INDEX NAME)

REFERENCE COUNT: 'AVAILABLE FOR THIS

44 THERE ARE 44 CITED REFERENCES

RE FORMAT

L12 ANSWER 10 OF 10 HCAPLUS COPYRIGHT 2006 ACS on STN

1981:157828 . HCAPLUS ACCESSION NUMBER:

94:157828 DOCUMENT NUMBER:

Flameproofing agent and flame-retardant TITLE:

plastic resin

compositions

Saito, Toranosuke; Ohishi, Hiroyuki INVENTOR(S):

Sanko Kaihatsu Kagaku Kenkyusho, Japan PATENT ASSIGNEE(S):

Ger. Offen., 36 pp. SOURCE:

CODEN: GWXXBX

DOCUMENT TYPE:

Patent

LANGUAGE:

German

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

	PATENT NO.	KIND	DATE	APPLICATION NO.
DATE				
-				
	 DE 2010275	7.1	19801002	DE 1980-3010375
	DE 3010375	A1 .	19801002	DE 1980-3010375
198003		CO	10041120	
	DE 3010375	C2	19841129	TD: 1070 01100
	JP 55124792	A2	19800926	JP 1979-31162
197903		- 4	10041004	
	JP 59053296	В4	19841224	
	JP _. 56104949	A2	19810821	JP 1980-6906
198001		,	•	
	FR 2451937	A1	19801017	FR 1980-5922
198003	317			
	FR 2451937	В1	19830909	
F	BE 882283	A1	19800918	BE 1980-199839
198003	318			•
1	NL 8001591	A	19800923.	NL 1980-1591 .
198003	318			
1	NL 186961	В	19901116	•
. 1	NL 186961	С	19910416	•
(GB 2049696	A	19801231	GB 1980-9104
198003	318			
(GB 2049696	В2	19830615	•
	JS 4317769	A	19820302	US 1980-131722
198003				
	ITY APPLN. INFO.:		•	JP 1979-31162 A
197903				
				JP 1980-6906 A

19800125

Alkali metal or alkaline earth salts of cyclic esters of (2'-hydroxy-2-

biphenylyl) phosphonic acid or its derivs. are flame retardants for

plastics. Thus, heating HOCH2CH2OH solns. of (9,10-dihydro-1,3-dichloro-9phospha-10-oxaphenanthren-9-oxide [61910-28-9] with NaOH at 170-208° for 8 h and acidification gives the corresponding cyclic phosphonate [76965-46-3], neutralization of which gives the Na salt (I) [76965-47-4]. Bisphenol A polycarbonate [24936-68-3] containing 2 phr I has UL-94 flammability rating V-O, compared with HB with no retardant. 61910-28-9 72741-96-9 72741-98-1 76964-81-3 76964-83-5 76964-85-7 76964-96-0 RL: RCT (Reactant); RACT (Reactant or reagent) (oxidation of) RN 61910-28-9 HCAPLUS 6H-Dibenz[c,e][1,2]oxaphosphorin, 2,4-dichloro-, 6-oxide (9CI) CN (CA INDEX NAME)

RN 72741-96-9 HCAPLUS
CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 2,4-dibromo-, 6-oxide (9CI)
(CA INDEX
NAME)

RN 72741-98-1 HCAPLUS
CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 2,4,8-trichloro-, 6-oxide
(9CI) (CA
INDEX NAME)

RN 76964-81-3 HCAPLUS
CN Ethanone, 1-(6-oxido-6H-dibenz[c,e][1,2]oxaphosphorin-2-yl)(9CI) (CA
INDEX NAME)

RN 76964-83-5 HCAPLUS
CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 4-chloro-, 6-oxide (9CI) (CA INDEX
NAME)

RN 76964-85-7 HCAPLUS
CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 2-chloro-, 6-oxide (9CI) (CA INDEX
NAME)

RN 76964-96-0 HCAPLUS
CN 6H-Dibenz[c,e][1,2]oxaphosphorin,
2,8-dichloro-4-(4-chlorophenyl)-,
6-oxide (9CI) (CA INDEX NAME)

RN 76964-82-4 HCAPLUS
CN Ethanone,
1-(6-hydroxy-6-oxido-6H-dibenz[c,e][1,2]oxaphosphorin-2-yl)(9CI) (CA INDEX NAME)

RN 76964-84-6 HCAPLUS
CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 4-chloro-6-hydroxy-, 6-oxide
(9CI) (CA
INDEX NAME)

RN 76964-86-8 HCAPLUS
CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 2-chloro-6-hydroxy-, 6-oxide
(9CI) (CA
INDEX NAME)

RN 76964-88-0 HCAPLUS CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 9-bromo-6-hydroxy-2,4,8-trimethyl-,

6-oxide (9CI) (CA INDEX NAME)

RN 76964-89-1 HCAPLUS
CN 6H-Dibenz[c,e][1,2]oxaphosphorin,
3,9-dibromo-6-hydroxy-2,4,8-trimethyl-,
6-oxide (9CI) (CA INDEX NAME)

RN 76964-91-5 HCAPLUS CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 9-fluoro-6-hydroxy-, 6-oxide (9CI) (CA INDEX NAME)

RN 76964-93-7 HCAPLUS
CN 6H-Dibenz[c,e][1,2]oxaphosphorin,
6-hydroxy-9-(trifluoromethyl)-, 6-oxide
(9CI) (CA INDEX NAME)

RN 76964-94-8 HCAPLUS
CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 2,4,8-trichloro-6-hydroxy-,6-oxide
(9CI) (CA INDEX NAME)

RN 76964-95-9 HCAPLUS CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 4-chloro-2-(1,1-dimethylethyl)-6-hydroxy-

, 6-oxide (9CI) (CA INDEX NAME)

RN 76965-46-3 HCAPLUS
CN 6H-Dibenz[c,e][1,2]oxaphosphorin, 2,4-dichloro-6-hydroxy-,
6-oxide (9CI)
(CA INDEX NAME)